AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF THE CLAIMS

Claims 1-43 (Cancelled).

- 44. (New) A method for reducing the glucoamylase activity in a milk clotting composition comprising the steps of:
- (i) providing a medium having a pH of 2.0 or higher that comprises chymosin activity and glucoamylase activity, wherein the medium having a pH of 2.0 or higher is derived from the cultivation of an organism that is a bacterial species, a yeast species, or a species of filamentous fungi, wherein the organism comprises a gene for encoding chymosin that is derived from a bovine or Camelidae species,
- (ii) lowering the pH of said medium having a pH of 2.0 or higher to between 1.0 and 1.8 by addition of lactic acid, acetic acid, propionic acid, or citric acid, and
- (iii) subjecting said medium to a pH in the range of 1.0 to 1.8 for a period of time sufficient to inactivate at least 50% of said glucoamylase activity while maintaining at least 75% of said chymosin activity.
- (New) The method according to claim 44, wherein at least 90% of said glucoamylase activity is inactivated.
- 46. (New) The method according to claim 44, wherein the medium having a pH of 2.0 or higher is a medium derived from the cultivation of an organism that during its cultivation produces said chymosin activity and said glucoamylase activity.
- 47. (New) The method according to claim 44, wherein the bacterial species is a gram negative bacterial species or a gram positive species.
- 48. (New) The method according to claim 46, wherein the bacterial species is E. coli or Racillus

- 49. (New) The method according to claim 44, where the yeast species is Saccharomyces cerevisiae, a methylotrophic yeast species or a Klyuveromyces species.
- 50. (New) The method according to claim 44, wherein the species of filamentous fungi is an Aspergillus species, a Cryphonectria species, a Fusarium species, a Rhizomucor species or a Trichoderma species.
- 51. (New) The method of claim 49, wherein said Aspergillus species is Aspergillus niger var.
- 52. (New) The method according to claim 44, wherein the medium having a pH of 2.0 or higher is subjected to a pH in the range of 1.5 to 1.8.
- 53. (New) The method according to claim 44, wherein the medium having a pH of 2.0 or higher is subjected to a pH between 1.7 to 1.8.
- 54. (New) The method according to claim 44, wherein the medium having a pH of 2.0 or higher is subjected to a pH of approximately 1.8.
- 55. (New) The method according to claim 44, wherein said period of time is in the range of 0.1 minutes to 48 hours.
- (New) The method according to claim 44, wherein the yeast species is selected from Pichia pastoris and Klyweromyces lactis.
- 57. (New) The method of claim 44, wherein the gene encoding chymosin is derived from Camelus dromedarius.
- 58. (New) The method of claim 44, wherein at least 85% of the chymosin activity is maintained in step (iii).
- 59. (New) The method of claim 44, wherein the gene encoding chymosin is derived from a bovine species.
- 60. (New) A method for reducing the glucoamylase activity in a milk clotting composition comprising the steps of:

- (i) providing a medium having a pH of 2.0 or higher that comprises chymosin activity and glucoamylase activity, wherein the medium having a pH of 2.0 or higher is derived from the cultivation of an organism that is a bacterial species, a yeast species, or a species of filamentous fungi, wherein the organism comprises a gene for encoding chymosin that is derived from a bovine or Camelidae species,
- (ii) lowering the pH of said medium having a pH of 2.0 or higher to between 1.0 and 1.8 by addition of an inorganic acid, and
- (iii) subjecting said medium to a pH in the range of 1.0 to 1.8 for a period of time sufficient to inactivate at least 50% of said glucoamylase activity while maintaining at least 75% of said chymosin activity.
- (New) The method according to claim 60, wherein at least 90% of said glucoamylase activity is inactivated.
- 62. (New) The method according to claim 60, wherein the medium having a pH of 2.0 or higher is a medium derived from the cultivation of an organism that during its cultivation produces said chymosin activity and said glucoamylase activity.
- 63. (New) The method according to claim 60, wherein the bacterial species is a gram negative bacterial species or a gram positive species.
- 64. (New) The method according to claim 63, wherein the bacterial species is E. coli or Racillus
- 65. (New) The method according to claim 60, where the yeast species is Saccharomyces cerevisiae, a methylotrophic yeast species or a Klywveromyces species.
- 66. (New) The method according to claim 60, wherein the species of filamentous fungi is an Aspergillus species, a Cryphonectria species, a Fusarium species, a Rhizomucor species or a Trichoderma species.
- (New) The method of claim 66, wherein said Aspergillus species is Aspergillus niger var.
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- 68. (New) The method according to claim 60, wherein the medium having a pH of 2.0 or higher is subjected to a pH in the range of 1.5 to 1.8.
- (New) The method according to claim 60, wherein the medium having a pH of 2.0 or higher is subjected to a pH between 1.7 to 1.8.
- 70. (New) The method according to claim 60, wherein the medium having a pH of 2.0 or higher is subjected to a pH of approximately 1.8.
- 71. (New) The method according to claim 60, wherein said period of time is in the range of 0.1 minutes to 48 hours.
- (New) The method according to claim 60, wherein the yeast species is selected from Pichia pastoris and Klyuveromyces lactis.
- 73. (New) The method of claim 60, wherein the gene encoding chymosin is derived from Camelus dromedarius.
- 74. (New) The method of claim 60, wherein at least 85% of the chymosin activity is maintained in step (iii).
- 75. (New) The method of claim 60, wherein the gene encoding chymosin is derived from a bovine species.
- (New) The method of claim 60, wherein the inorganic acid is hydrochloric acid, phosphoric acid, or sulfuric acid.
- 77. (New) The method of claim 60, wherein the gene encoding chymosin is derived from Camelus dromedarius.